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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/599,451	07/18/2007	Domenico Fanara	06-796	9142
20306	7590	01/05/2010		
MCDONNELL BOEHNEN HULBERT & BERGHOFF LLP			EXAMINER	
300 S. WACKER DRIVE			THOMAS, TIMOTHY P	
32ND FLOOR			ART UNIT	PAPER NUMBER
CHICAGO, IL 60606			1628	
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			01/05/2010	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Examiner-Initiated Interview Summary	Application No.	Applicant(s)	
	10/599,451	FANARA ET AL.	
	Examiner TIMOTHY P. THOMAS	Art Unit 1628	

All Participants: **Status of Application:** _____

(1) TIMOTHY P. THOMAS. (3) _____.

(2) Michael Greenfield. (4) _____.

Date of Interview: 30 December 2009

Time: 9:00 a.m.

Type of Interview:

Telephonic
 Video Conference
 Personal (Copy given to: Applicant Applicant's representative)

Exhibit Shown or Demonstrated: Yes No

If Yes, provide a brief description: _____.

Part I.

Rejection(s) discussed:

See Continuation Sheet

Claims discussed:

1, 6-8, 14-15, 18, 22

Prior art documents discussed:

Doron et al. ("Antibacterial effect of parabens against planktonic and biofilm *Streptococcus sobrinus*," 2001 *International Journal of Antimicrobial Agents*; 18: 575-578)

Part II.

SUBSTANCE OF INTERVIEW DESCRIBING THE GENERAL NATURE OF WHAT WAS DISCUSSED:

See Continuation Sheet

Part III.

It is not necessary for applicant to provide a separate record of the substance of the interview, since the interview directly resulted in the allowance of the application. The examiner will provide a written summary of the substance of the interview in the Notice of Allowability.
 It is not necessary for applicant to provide a separate record of the substance of the interview, since the interview did not result in resolution of all issues. A brief summary by the examiner appears in Part II above.

/Timothy P Thomas/
Examiner, Art Unit 1628

(Applicant/Applicant's Representative Signature – if appropriate)

Continuation of rejections discussed: Claims 1-2, 5, 12 and 17, rejected under 35 U.S.C. 103(a) as being unpatentable over DeLongueville et al. (WO 02/47689 A2; IDS 12/18/2008 reference); Gilliland et al. (Gilliland 1) ("The bactericidal activity of a methyl and propyl parabens combination: isothermal and non-isothermal studies"; 1992; Journal of Applied Bacteriology; 72: 252-257); Gilliland et al. (Gilliland 2) ("Kinetic evaluation of claimed synergistic paraben combinations using a factorial design"; 1992; Journal of Applied Bacteriology; 72: 258-261); in view of Routledge et al. ("Some Alkyl Hydroxy Benzoate Preservatives (Parabens) Are Estrogenic"; 1998; Toxicology and Applied Pharmacology; 153: 12-19).

Continuation of Substance of Interview including description of the general nature of what was discussed: The reply of record is considered to be persuasive with respect to the prior art rejection of record, based on an unexpected result disclosed in the specification. However, the upper limit of claim 1 has been increased to 1.125 mg/ml. Looking at the Doron reference, which is of record, the reference is suggestive that an amount of just above 0.9% [MP] + [PP] (0.06 + 0.03) would be expected to result in solutions with 0% bacterial growth, based on the data of Figure 1; i.e., 0% bacterial growth would be expected from interpolation of the data at concentrations of just under 1.125 mg/mL. It was proposed that the claim be limited to 1 mg/mL or less as the upper amount, in place of "less than 1.125 mg/mL". Since the basis of overcoming a rejection that would include the Doron reference would be unexpected results, the lowest amount for which there is an unexpected result in the specification is 0.375 mg/mL. It was therefore proposed that the lower limit of claim 1 be 0.375 or greater, in place of the amount of more than 0, currently in claim 1. The range of 0.375 to 1 mg/mL is commensurate in scope to the data for which unexpected results are present in the disclosure.

It was noted that should claim 1 be amended as proposed, and the dependent claims be amended in a corresponding manner, the case would be discussed with supervisors for consideration of whether the claims are patentable. Considering the withdrawn claim components of claims 6-8, each of these compounds are known antimicrobial agents; inclusion of these compounds would no longer lead to an unexpected result when any of the additional compounds are present; i.e., should claim 1 be determined to be allowable, claims 6-8 are not automatically also allowable. Claim 18 would need to contain the limitations of claim 1 to be considered for rejoinder; as would claim 22, should claim 1 be determined to be allowable. .